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Clinic: Columbia

Date: 3/15/2017

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Public	Departments\Clinic Forms\STAFF EDUCATION-REEDUCATION FORM 9-2012



Missouri Departin. Health and Senior Services P.O. Box 570, Jefferson City, MO 65102-0570 Phone: 573-751-6400 FAX: 573-751-6010 RELAY MISSOURI for Hearing and Speech Impaired 1-800-735-2966 VOICE 1-800-735-2466 Randall W. Williams, MD, FACOG Director



Eric R. Greitens Governor

October 3, 2017

Vicki Casey Comprehensive Health of Planned Parenthood Great Plains, Inc. 711 North Providence Road Columbia, MO 65203

Re: Comprehensive Health of Planned Parenthood Great Plains, Inc. - Columbia survey

Dear Ms. Casey:

Please see attached results of the recent follow-up survey of August 28, 2017. Your facility is now in compliance with current legal requirements for licensure.

Please retain this material for your own records. The abortion facility license is attached, effective date October 3, 2017.

We welcome any questions at 573-751-6083.

Respectfully,

John Farystu

John Langston, Administrator John.Langston@health.mo.gov Bureau of Ambulatory Care Missouri Department of Health & Senior Services

www.health.mo.gov

Healthy Missourians for life. The Missouri Department of Health and Senior Services will be the leader in promoting, protecting and partnering for health.

ND PLAN	NT OF DEFICIENCIES	Alth and Senior Services (X1) PROVIDER/SUPPLIER/CLIA IDENTIFICATION NUMBER:	(X2) MULTIPLE CONSTRUCTION		(X3) DAT	(X3) DATE SURVEY	
			A. BUILDING:			COMPLETED	
		A004	B. WING			R 08/28/2017	
AME OF	PROVIDER OR SUPPLIER	STREET A	DDRESS, CITY, S	TATE, ZIP CODE		20/2011	
OMPRI	EHENSIVE HEALTH F		ROVIDENCE R BIA, MO 6520:				
(X4) ID PREFIX		ATEMENT OF DEFICIENCIES	ID	PROVIDER'S PLAN OF	CORRECTION	(X5	
TAG	(EACH DEFICIENCY MUST BE PRECEDED BY FULL REGULATORY OR LSC IDENTIFYING INFORMATION)		PREFIX TAG	(EACH CORRECTIVE AC CROSS-REFERENCED TO DEFICIENC	TION SHOULD BE		
{L 000}	Initial Comments		{L 000}				
	An onsite Licensure revisit survey was conducted on 08/28/17. The facility was found to be in substantial compliance with the rules and regulations for abortion facilities found at 19 CSR 30-30.060.						
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Randall W. Williams, MD, FACOG Director



Eric R. Greitens Governor

August 11, 2017

Amanda Addison (<u>Amanda.addison@ppgreatplains.org</u>) Comprehensive Health of Planned Parenthood Great Plains 711 North Providence Road Columbia, Mo 65203

Re: Comprehensive Health of Planned Parenthood Great Plains - Columbia Revisit Survey

Dear Ms. Addison:

The Department received the application for licensure of the Columbia Planned Parenthood location as an abortion facility. Department staff conducted an onsite survey of the facility on October 11, 2016 to determine compliance with the terms of the 2010 settlement agreement and applicable statutes and regulations. As a result, on November 2, 2016, your facility was provided with a list of regulatory items that were not in compliance.

After the facility submitted a complete response and documentation regarding correction of the items that were not in compliance, the Department performed an onsite revisit of the facility on July 25, 2017.

Listed below are items the revisit survey indicated were still not in compliance. Until a written response is provided describing how all items below have been addressed, including acceptable evidence of compliance, an abortion facility license cannot be issued.

19 CSR 30-30.0601(B)8. The facility shall establish a program for identifying and preventing infections and for maintaining a safe environment.

The facility failed to demonstrate compliance with facility's established Infection Program, based on Association for the Advancement of Medical Instrumentation (AAMI) and Centers for Disease Control (CDC) standards. Specific findings:

1. The facility failed to follow the manufacturer's instructions for use (IFU) for routine care of the sterilizers.

(AAMI 9.4 Routine Care: Sterilizers should be inspected and cleaned daily according to the manufacturers' written instructions. Weekly or other prescribed inspection and cleaning should be performed as specified in the manufacturers' written IFU.)

2. The facility failed to maintain a separate autoclave log with the required components tracked for each of two sterilizers.

(AAMI 10.3.2 Sterilizer records: For each sterilization cycle, the following information should be recorded and maintained: Lot number; Specific contents of the lot or load, including quantity, department, and a specific description of the items [e.g. towels, type/name of instrument sets]; Exposure time and temperature, if not provided on the sterilizer recording chart; Name or initials of the operator; Results of biological testing, if applicable; Any reports of inconclusive or nonresponsive chemical indicators found later in the load.)

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Healthy Missourians for life. The Missouri Department of Health and Senior Services will be the leader in promoting, protecting and partnering for health. 3. The facility failed to maintain a record of sterilizer maintenance and repair. (AAMI 9.7 A maintenance record, in either paper or electronic format, should be kept for each sterilizer. At

least the following information should be recorded:

- *a) The date on which service was requested;*
- *b)* The model and serial number of the sterilizer;
- *c) The location of the equipment (if applicable);*
- *d) The name of the individual from the facility who requested and authorized the service;*
- *e)* The reason for the service request;
- f) A description of the service performed;
- g) The types an quantities of parts replaced;
- *h)* The name of the person who performed the service;
- *i) The date the work was completed;*
- *j) The handwritten or electronic signature and title of the person who acknowledged completion of the work; and*
- *k)* The results of any post-maintenance testing performed, if needed, before the sterilizer was returned to service.)
- 4. The facility failed to have written processes for reprocessing and/or quarantine of instruments following positive biological indicators, as well as consecutive biological testing following sterilizer failure and repair.
- 5. During an interview with facility advanced practice nursing staff at the time of the revisit, staff acknowledged the problems with adequate documentation and clear adherence to the AAMI standards.

Please respond in writing providing evidence/documentation that each of these items has been fully addressed and corrected.

If you have further questions, you may contact our office at 573-751-6083 or via email at the address noted below.

Sincerely,

John Jarysta

John Langston, Administrator John.Langston@health.mo.gov Bureau of Ambulatory Care Missouri Department of Health & Senior Services